

Product datasheet

Anti-PIP5KI gamma antibody [MAO-R1] ab109192

**重组** RabMAb

2 References 1 图像

概述

<b>产品名称</b>	Anti-PIP5KI gamma抗体[MAO-R1]
<b>描述</b>	兔单克隆抗体[MAO-R1] to PIP5KI gamma
<b>宿主</b>	Rabbit
<b>经测试应用</b>	<b>适用于:</b> WB, IP <b>不适用于:</b> Flow Cyt, ICC or IHC-P
<b>种属反应性</b>	<b>与反应:</b> Mouse, Rat, Human
<b>免疫原</b>	Recombinant full length protein within Mouse PIP5KI gamma. The exact sequence is proprietary.
<b>阳性对照</b>	K562, Human fetal kidney, C6, Neuro-2a, and 293T cell lysates
<b>常规说明</b>	

Our RabMAb<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to [RabMAb<sup>®</sup> patents](#)

This product is a recombinant rabbit monoclonal antibody.

性能

<b>形式</b>	Liquid
<b>存放说明</b>	Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.
<b>存储溶液</b>	pH: 7.20 Preservative: 0.05% Sodium azide Constituents: 0.1% BSA, 40% Glycerol, 9.85% Tris glycine, 50% Tissue culture supernatant
<b>纯度</b>	Tissue culture supernatant
<b>克隆</b>	单克隆
<b>克隆编号</b>	MAO-R1
<b>同种型</b>	IgG

应用

Our [Abpromise guarantee](#) covers the use of **ab109192** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

应用	Ab评论	说明
WB		1/1000 - 1/10000. Detects a band of approximately 80-90 kDa (predicted molecular weight: 73 kDa).
IP		1/10 - 1/100.

**应用说明** Is unsuitable for Flow Cyt, ICC or IHC-P.

## 靶标

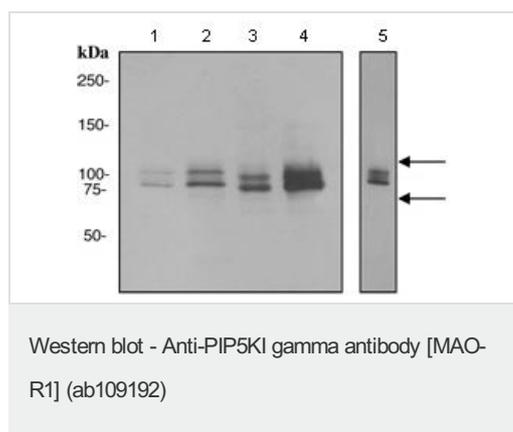
**功能** Plays a role in membrane ruffling and assembly of clathrin-coated pits at the synapse. Mediates RAC1-dependent reorganization of actin filaments (By similarity). Participates in the biosynthesis of phosphatidylinositol-4,5-bisphosphate.

**疾病相关** Defects in PIP5K1C are the cause of lethal congenital contracture syndrome type 3 (LCCS3) [MIM:611369]; also known as multiple contractural syndrome Israeli Bedouin type B. LCCS is an autosomal recessive disorder characterized by early fetal hydrops and akinesia, the Pena-Shokeir phenotype, specific neuropathology with degeneration of anterior horn neurons and extreme skeletal muscle atrophy. LCCS3 patients present at birth with severe multiple joint contractures with severe muscle wasting and atrophy, mainly in the legs. LCCS3 can be distinguished from the original LCCS by the absence of hydrops, fractures, and multiple pterygia.

**序列相似性** Contains 1 PIPK domain.

**细胞定位** Cell membrane. Cytoplasmic, associated with the plasma membrane. Detected in focal adhesion plaques, membrane ruffles and plasma membrane invaginations.

## 图片



**All lanes :** Anti-PIP5K1 gamma antibody [MAO-R1] (ab109192) at 1/1000 dilution

**Lane 1 :** K562 cell lysate

**Lane 2 :** Human fetal kidney cell lysate

**Lane 3 :** C6 cell lysate

**Lane 4 :** Neuro-2a cell lysate

**Lane 5 :** 293T cell lysate

Lysates/proteins at 10 µg per lane.

**Predicted band size:** 73 kDa

**Please note:** All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE"

## **Our Abpromise to you: Quality guaranteed and expert technical support**

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- Replacement or refund for products not performing as stated on the datasheet
- Valid for 12 months from date of delivery
- Response to your inquiry within 24 hours
  
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- We investigate all quality concerns to ensure our products perform to the highest standards

If the product does not perform as described on this datasheet, we will offer a refund or replacement. For full details of the Abpromise, please visit <https://www.abcam.cn/abpromise> or contact our technical team.

## **Terms and conditions**

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